## DEPARTMENT OF JUSTICE

**Drug Enforcement Administration** 

Bulk Manufacturer of Controlled Substances Application: AMRI Rensselaer, Inc.
[Docket No. DEA-392]

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.33(a) on or before [INSERT 60 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER].

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrissette Drive, Springfield, Virginia 22152.

## SUPPLEMENTARY INFORMATION:

The Attorney General has delegated his authority under the Controlled Substances

Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR

0.100(b). Authority to exercise all necessary functions with respect to the promulgation
and implementation of 21 CFR part 1301, incident to the registration of manufacturers,
distributors, dispensers, importers, and exporters of controlled substances (other than
final orders in connection with suspension, denial, or revocation of registration) has been
redelegated to the Assistant Administrator of the DEA Diversion Control Division

("Assistant Administrator") pursuant to section 7 of 28 CFR part 0, appendix to subpart

R.

In accordance with 21 CFR 1301.33(a), this is notice that on June 25, 2018, AMRI Rensselaer, Inc., 33 Riverside Avenue, Rensselaer, New York 12144 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled Substance	Drug Code	Schedule
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
Amphetamine	1100	II
Lisdexamfetamine	1205	II
Pentobarbital	2270	II
4-Anilino-N-phenethyl-4-piperidine (ANPP)	8333	II
Codeine	9050	II
Oxycodone	9143	II
Hydromorphone	9150	II
Hydrocodone	9193	II
Meperidine	9230	II
Morphine	9300	II

The company plans to manufacture bulk controlled substances for use in product development and for distribution to its customers.

In reference to drug codes 7360 (marihuana) and 7370 (THC), the company plans to bulk manufacture these drugs as synthetics. No other activities for these drug codes are authorized for this registration.

Dated: July 23, 2018.

John J. Martin,

Assistant Administrator.

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